



Murine Bone Marrow Mesenchymal Stromal Cells Respond Efficiently to Oxidative Stress Despite the Low Level of Heme Oxygenases 1 and 2

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Abstract

Aims: Mesenchymal stromal cells (MSCs) are heterogeneous cells from adult tissues that are able to differentiate *in vitro* into adipocytes, osteoblasts, or chondrocytes. Such cells are widely studied in regenerative medicine. However, the success of cellular therapy depends on the cell survival. Heme oxygenase-1 (HO-1, encoded by the *Hmox1* gene), an enzyme converting heme to biliverdin, carbon monoxide, and Fe²⁺, is cytoprotective and can affect stem cell performance. Therefore, our study aimed at assessing whether *Hmox1* is critical for survival and functions of murine bone marrow MSCs.

Results: Both MSC *Hmox1*^{+/+} and *Hmox1*^{-/-} showed similar phenotype, differentiation capacities, and production of cytokines or growth factors. *Hmox1*^{+/+} and *Hmox1*^{-/-} cells showed similar survival in response to 50 μmol/L hemin even in increased glucose concentration, conditions that were unfavorable for *Hmox1*^{-/-} bone marrow-derived proangiogenic cells (BDMC). *Hmox1*^{+/+} MSCs but not fibroblasts retained low ROS levels even after prolonged incubation with 50 μmol/L hemin, although both cell types have a comparable *Hmox1* expression and similarly increase its levels in response to hemin. MSCs *Hmox1*^{-/-} treated with hemin efficiently induced expression of a vast panel of antioxidant genes, especially enzymes of the glutathione pathway.

Innovation and Conclusion: *Hmox1* overexpression is a popular strategy to enhance viability and performance of MSCs after the transplantation. However, murine MSCs *Hmox1*^{-/-} do not differ from wild-type MSCs in phenotype and functions. MSC *Hmox1*^{-/-} show better resistance to hemin than fibroblasts and BDMCs and rapidly react to the stress by upregulation of quintessential genes in antioxidant response. *Antioxid. Redox Signal.* 29, 111–127.

Keywords: stem cells, antioxidant gene response, heme, mesenchymal stem cell

Introduction

MESENCHYMAL STROMAL CELLS (MSCs), also known as mesenchymal stem cells, or multipotent stromal cells are a heterogeneous population of connective tissue cells that contains osteoblast and adipocyte progenitors, fibroblasts, and smooth muscle cells (5). *In vitro* criteria for human MSCs

include adherence to the plastic in standard culture conditions, differentiation *in vitro* to adipocytes, osteoblasts, and chondrocytes (9). MSCs should express CD73, CD90, and CD105 markers but not CD45, CD34, CD14, CD11b, CD79α, CD19, and HLA-DR (9). MSCs were further identified, also *in vivo*, by CD271 and CD106 (human) (33), CD146 (44), nestin (34), Sca-1 (mouse) and PDGFRα

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Innovation

Enhancement of stem and progenitor cells antioxidant capacity is the aim of many studies focusing on the cellular therapies. Many of such strategies propose the overexpression of *Hmox1* as a protection against cell stress. For the first time, this article shows that mesenchymal stromal cells (MSCs) lacking *Hmox1* can more efficiently than other cells deal with oxidative stress induced with hemin, using the mechanism involving the upregulation of glutathione pathway. High resistance to stress and unique ability to activate antioxidant response suggest that MSC may not need additional protection by *Hmox1* overexpression.

(36), leptin receptor LepR (8, 67), or high expression of CXCL12 (41).

MSCs were shown to be immune evasive or immunomodulatory, depending on the microenvironment (2). The mechanism of immunosuppression is complex and involves many factors, that is, prostaglandin E₂, nitric oxide, and TGF β (39). Although MSCs are commonly believed to deal with oxidative stress efficiently (55), the biggest obstacle to the therapeutic use of MSCs is their poor survival and engraftment after the transplantation (11). Therefore, many studies focus on the enhancement of their antioxidant activity with overexpression of various genes, for example, *Hmox1* (54, 63).

Heme oxygenase-1 (HO-1, encoded by the *HMOX1* gene) is an enzyme degrading heme to carbon monoxide (CO), biliverdin, and Fe²⁺ ions. Due to its enzymatic activity, heme oxygenase-1 influences cell survival, resistance to the oxidative stress, and angiogenesis (10). We have recently shown that proangiogenic cells isolated from the bone marrow of *Hmox1* knock-out mice present impaired proliferation, migration, and formation of capillaries (16). What is more, overexpression of heme oxygenase-1 can lead to the block of differentiation, that is, in myoblasts (27).

Rat MSCs transfected with the plasmid coding for human heme oxygenase-1 showed decreased apoptosis in hypoxia and higher resistance to H₂O₂ (54). In our hands, pig bone marrow-derived cells transduced with adenoviral vectors encoding heme oxygenase-1 (AdHO1) were characterized by better angiogenic activity *in vitro* and improved left ventricular ejection fraction 30 min after infarction in pigs (63). Treatment with cobalt protoporphyrin IX (CoPP), heme oxygenase-1 activator, enhanced proliferation of human mesenchymal stem cells and production of VEGF; whereas tin protoporphyrin IX (SnPP), heme oxygenase-1 inhibitor, had an opposite influence (20). Further, CoPP-treated MSCs accelerated wound healing in a xenogeneic model of diabetic mice (20).

Modulation of heme oxygenase-1 activity with SnPP in human MSCs affected their ability to inhibit T cell proliferation *in vitro*. Interestingly, the effect of SnPP was not observed in rat MSCs, and T cell proliferation was restored only when concomitant treatment of nitric oxide synthase 2 was used (7). Moreover, heme oxygenase inhibition decreased the ability of MSCs to induce Tr1 and Th3 regulatory cells and to elevate levels of IL-10 and TGF β , respectively. MSCs preconditioned with a mixed lymphocyte reaction showed decreased HO-1 levels as well as immunomodulatory activity (37).

Finally, the effect of heme oxygenase-1 on MSC differentiation to adipocytes and osteoblasts was also studied. Abraham and co-workers showed in the series of publications that enhanced expression of HO-1 in MSCs results in improved differentiation to osteoblasts, whereas its inhibition promotes adipogenesis (3, 56–59). On the other hand, Zarjou *et al.* reported no differences in differentiation potential between MSC *Hmox1*^{+/+} and *Hmox1*^{-/-} (66). Also in other studies, overexpression of heme oxygenase-1 in MSCs did not affect their differentiation (18, 68).

Data on the influence of heme oxygenase-1 on MSCs are often contradictory. Conjointly, copper or tin protoporphyrins were used in many studies to modulate HO-1 activity, although they were shown to have many heme oxygenase-independent effects in various cell types (17, 23). MSCs are essential for the proper function of stem cell niches in bone marrow, and lack of heme oxygenase-1 was shown to potentially affect other bone marrow-derived cells, that is, pro-angiogenic cells (PACs) (16). Therefore, we decided to characterize murine bone marrow-derived MSCs lacking the functional *Hmox1* gene, with the focus on their response to oxidative stress.

Results

Hmox1^{+/+} or *Hmox1*^{-/-} bone marrow MSCs show similar phenotype and differentiation

First, we compared the phenotypes of murine bone marrow stromal cells *Hmox1*^{+/+} or *Hmox1*^{-/-} in culture by using flow cytometry. Regardless of the genotype, 60% of the cells in culture were CD45⁻CD31⁻ (data not shown). Therefore, cells used for all the experiments were purified from the remaining CD45⁺ fraction with MACS sorting. Obtained cells lacked expression of endothelial markers CD31 and CD34, whereas CD117 (c-kit) was expressed only on a small sub-fraction of cells (Fig. 1A). Isolated MSCs expressed positive markers that were attributed to the mesenchymal stem/stromal cells, that is, CD29, CD90, CD105, Ly-6A/E (Sca-1) (Fig. 1B), and CD140a (PDGFR α) (Fig. 1C, D). *Hmox1*^{-/-} bone marrow-derived PACs had impaired proliferation (16). However, the proliferation of MSC *Hmox1*^{+/+} or *Hmox1*^{-/-} was similar, even when cells were grown under stress conditions in high glucose concentration (Fig. 2A).

Further, bone marrow stromal cells isolated from long bones of mice *Hmox1*^{+/+} or *Hmox1*^{-/-} formed similar numbers of colonies that consisted of fibroblastoid cells (Fig. 2B) or cells able to be differentiated to osteoblasts (Fig. 2C). MSCs were shown to differentiate into adipocytes, osteoblasts, and chondrocytes (6). Both *Hmox1*^{+/+} and *Hmox1*^{-/-} MSCs differentiated into osteoblasts and adipocytes, which were evidenced with staining for osteopontin or Fabp4, respectively (Supplementary Fig. S1A, B; Supplementary Data are available online at www.liebertpub.com/ars).

Because HO-1 was suggested to play a crucial role in the regulation of MSC adipogenesis (59), we focused on the effects of *Hmox1* knockout on the genes associated with lipid metabolism. Interestingly, differentiation of murine bone marrow MSCs to adipocytes induced similar changes in the gene expression in both *Hmox1*^{+/+} and *Hmox1*^{-/-} cells (Supplementary Fig. S1C).

However, *Hmox1*^{+/+} MSCs were the only ones that increased expression of miR-21-5p, the microRNA that, via TGF β signaling, regulates adipogenesis (26) (Supplementary

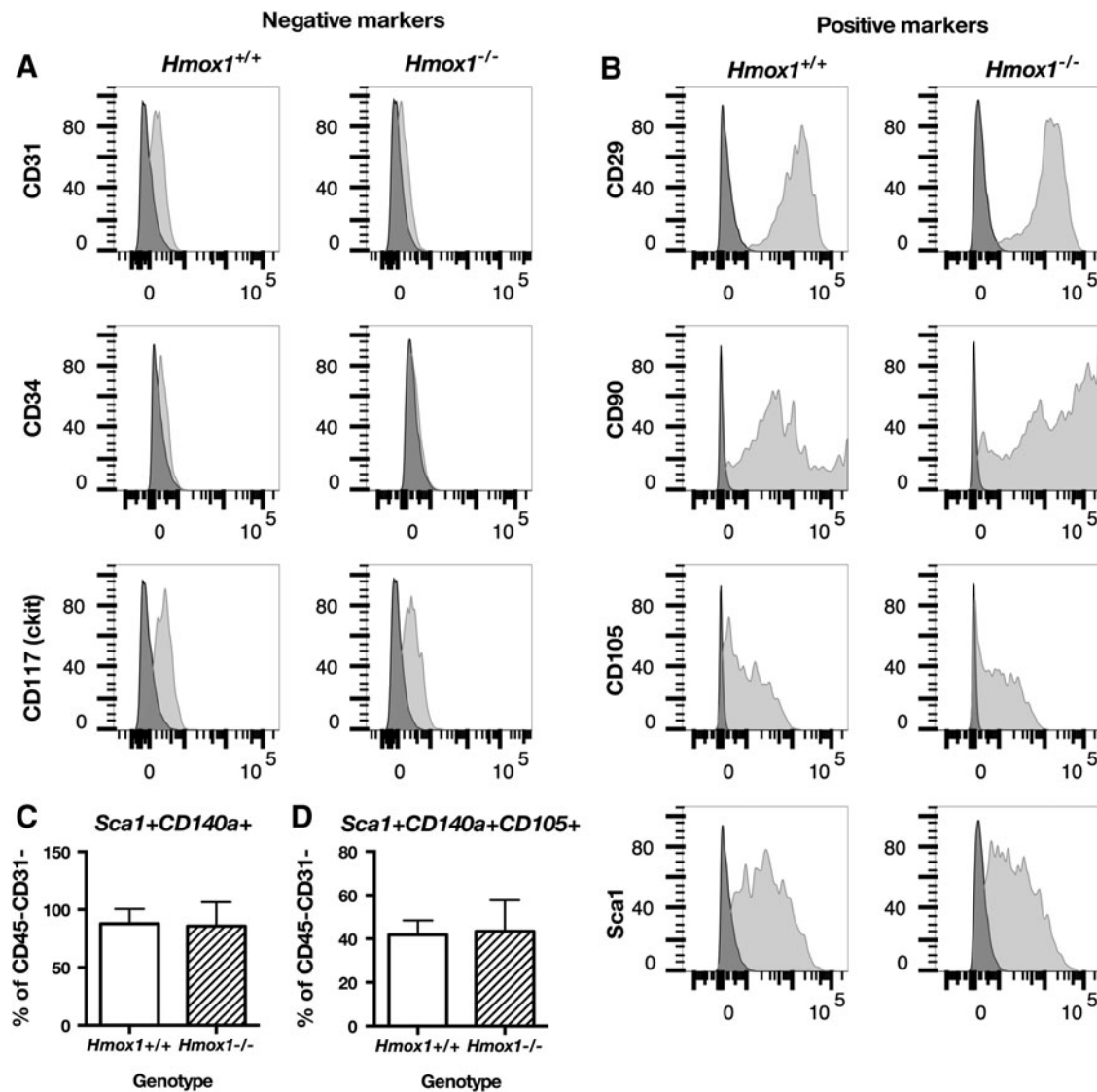


FIG. 1. MSC *Hmox1*^{+/+} or *Hmox1*^{-/-} have similar phenotype. Expression of MSC positive markers CD29, CD90, CD105, and Sca-1 (A), MSC negative markers CD31, CD34, and ckit in *Hmox1*^{+/+} or *Hmox1*^{-/-} MSCs (B). Phenotype of nonsorted bone marrow stromal cells in passage 4: fraction of CD45⁻CD31⁻ cells in culture (C), fraction of Sca-1⁺CD140a⁺ cells within CD45⁻CD31⁻ population in *Hmox1*^{+/+} or *Hmox1*^{-/-} cells. (C, D, N=9–10). MSCs, mesenchymal stromal cells.

Fig. S1D). Levels of other tested microRNAs associated with adipogenesis, namely miR-31-5p, miR-150-5p, miR-301a-5p, miR-378a-3p, or miR-378a-5p, remained unchanged in cells of both genotypes (Supplementary Fig. S1E–I). Further, basal expression of all tested microRNAs, including miR-21-5p, which increased during adipocyte differentiation only in *Hmox1*^{+/+} cells, was similar in MSC *Hmox1*^{+/+} and *Hmox1*^{-/-} (data not shown).

Effects of changed expression of *HMOX1* in human MSCs on the adipogenesis or osteogenesis were stronger when cells were cultured in high glucose concentration (3). Therefore, we tested markers of adipogenesis in MSC *Hmox1*^{+/+} or *Hmox1*^{-/-} differentiated in low or high glucose concentration. Interestingly, adipogenic differentiation of murine bone marrow-derived *Hmox1*^{+/+} or *Hmox1*^{-/-} MSCs did not change when cells were cultured under high glucose conditions. Both *Hmox1*^{+/+} and *Hmox1*^{-/-} MSCs upregulated

fatty acid-binding protein 4—a marker of adipocyte differentiation (Fig. 2D).

MSCs were shown to be precursors of fibroblasts and myofibroblasts and, therefore, contribute to the tumor stroma (35) or development of fibrosis (30). Heme oxygenase-1 can affect both tumor microenvironment (62) and kidney fibrosis (47). Therefore, we investigated whether the lack of *Hmox1* gene in MSCs may influence their ability to form myofibroblasts. *Hmox1*^{+/+} and *Hmox1*^{-/-} MSCs were differentiated to myofibroblasts with TGFβ1 treatment for 6 days. Cells changed their morphology (Supplementary Fig. S1J) and upregulated α-smooth muscle actin (*Acta2*) (Fig. 2E). Of note, upregulation of *Acta2* was lower in myofibroblasts derived from *Hmox1*^{-/-} cells.

Transcript levels of fibroblast-specific protein 1 (*Fsp1*) (Fig. 2F) tended to decrease in *Hmox1*^{+/+} and *Hmox1*^{-/-} MSCs; however, this trend reached significance only in

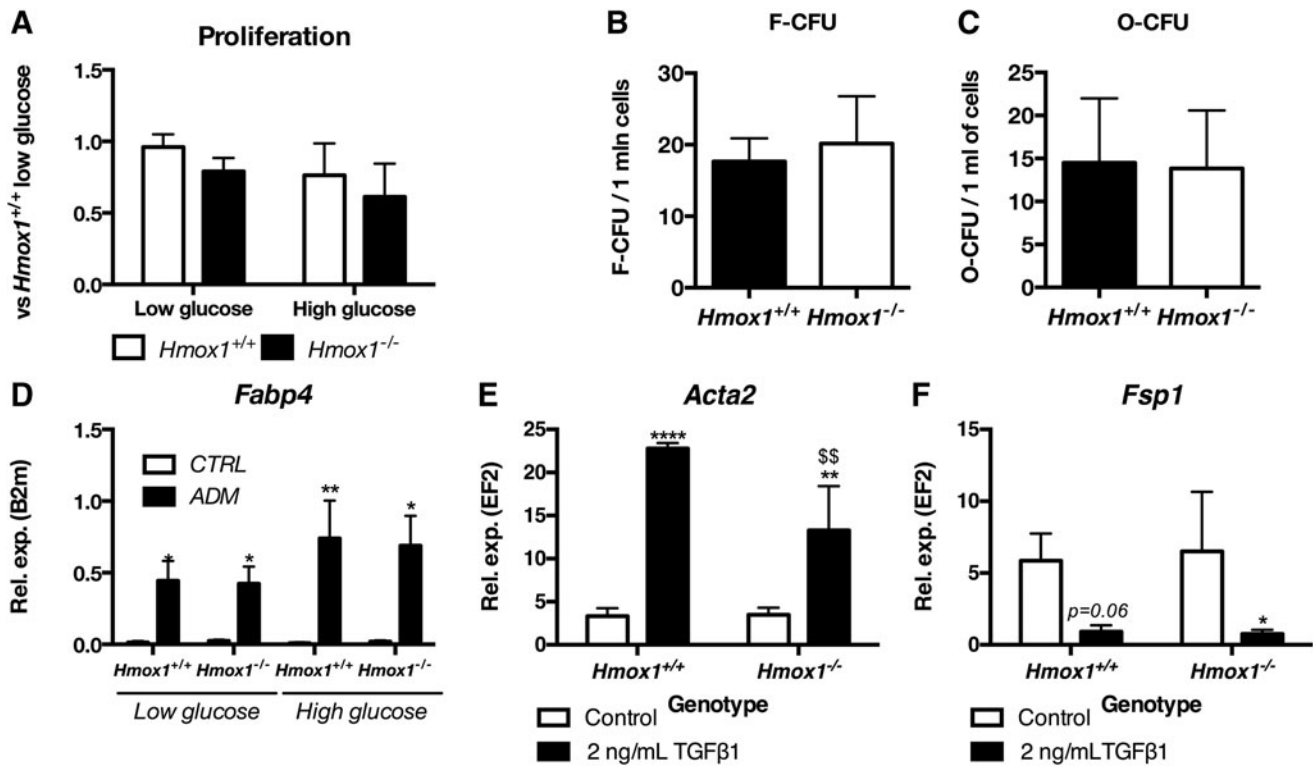


FIG. 2. MSC $Hmox1^{+/+}$ or $Hmox1^{-/-}$ show similar ability to differentiate to adipocytes, osteoblasts and smooth muscle cells. Proliferation of MSC $Hmox1^{+/+}$ or $Hmox1^{-/-}$ in low or high glucose medium assessed with BrDU assay (A). Ability to form fibroblastoid (B) or osteoblast (C) colonies by bone marrow cells isolated from $Hmox1^{+/+}$ or $Hmox1^{-/-}$ MSCs assessed with fibroblastoid or osteoblast colony-forming unit assay, respectively. Data shown as mean \pm SD, $N=3$. Expression of *Fabp4* in $Hmox1^{+/+}$ or $Hmox1^{-/-}$ MSCs differentiated to adipocytes in low or high glucose conditions (D). Gene expression was assessed with qRT-PCR. Data are shown as mean \pm SD, $N=4-7$. Expression of *Acta2* (E), *Fsp1* (F) measured with qRT-PCR in control and differentiated MSC $Hmox1^{+/+}$ or $Hmox1^{-/-}$. Data are shown as mean \pm SD, $*p<0.05$, $**p<0.01$, $****p<0.0001$ control versus TGF β 1; $^{SS}p<0.01$ $Hmox1^{+/+}$ versus $Hmox1^{-/-}$, $N=3$. qRT-PCR, quantitative real-time PCR.

$Hmox1^{-/-}$ cells. To summarize, we show here that MSCs, regardless of the heme oxygenase-1, show similar phenotype and differentiation ability, even in stress conditions.

Lack of *Hmox1* does not change MSCs immunomodulatory activity or production of cytokines

Akiyama *et al.* showed that MSCs injected intravenously can decrease numbers of circulating CD3 $^{+}$ and increase T cell apoptosis (1). Mougiakakos *et al.* suggested that inhibition of HO-1 activity can decrease MSCs ability to induce T regulatory cells (37). Therefore, to test the effect of complete knock-out of *Hmox1* on immunomodulatory activity of MSCs, we injected wild-type C57Bl6 \times FVB mice with MSC $Hmox1^{+/+}$ or $Hmox1^{-/-}$ and analyzed apoptosis and activation of T cells.

Numbers of circulating CD3 $^{+}$ T cells did not change in mice injected with MSCs (Fig. 3A). However, numbers of apoptotic Annexin V $^{+}$ CD3 $^{+}$ cells tended to increase after the MSC injection, but this trend did not reach statistical significance (Fig. 3B). Injection of MSC $Hmox1^{+/+}$ or $Hmox1^{-/-}$ did not affect the numbers as well of activated CD3 $^{+}$ CD4 $^{+}$ CD25 high T cells (Fig. 3C). *In vitro*, primary murine CD3 $^{+}$ T cells co-cultured with MSC $Hmox1^{+/+}$ or $Hmox1^{-/-}$ showed a similar cell cycle (Supplementary Fig. S2A–C), percentage of Ki67 $^{+}$ cells (Supplementary Fig. S2D) and annexin V $^{+}$ cells (Supplementary Fig. S2E).

Further, lack of the functional *Hmox1* gene did not change the profile of cytokines and growth factors produced by MSC. Conditioned media from MSC $Hmox1^{+/+}$ and $Hmox1^{-/-}$ contained similar amounts of G-CSF, IL-6, LIF, LIX, CXCL1, CXCL2, CCL5, and VEGF (Fig. 3D–K). Levels of eotaxin, GM-CSF, IG γ , IL-1 α , IL-1 β , IL-2, IL-3, IL-4, IL-5, IL-7, IL-9, IL-10, IL-12 p40, IL-12 p70, IL-13, IL-15, IL-17, MIP-1 α , MIP-1 β , M-CSF, MIP-2, MIG, and TNF α were under the threshold of detection. Therefore, we concluded that lack of the *Hmox1* gene did not change the tested MSC secretome and did not affect the ability of MSCs to induce T cell death both *in vitro* and *in vivo*.

$Hmox1^{+/+}$ and $Hmox1^{-/-}$ MSCs show high resistance to oxidative stress induced with H $_2$ O $_2$ or hemin

Bone marrow-derived proangiogenic cells (BDMC) lacking *Hmox1* gene were characterized by higher sensitivity to oxidative stress, that is, induced with hemin (16). Therefore, we decided to test whether it is also true for MSCs. Surprisingly, $Hmox1^{+/+}$ and $Hmox1^{-/-}$ MSCs showed no difference in the viability when treated for 6 h with H $_2$ O $_2$ or 50 μ mol/L hemin in both low and high glucose conditions (Fig. 4A, B). Moreover, treatment with 10 ng/ μ L of TNF α for 24 h did not yield significant differences in the viability of MSCs with or without *Hmox1* (Supplementary Fig. S3A, B).

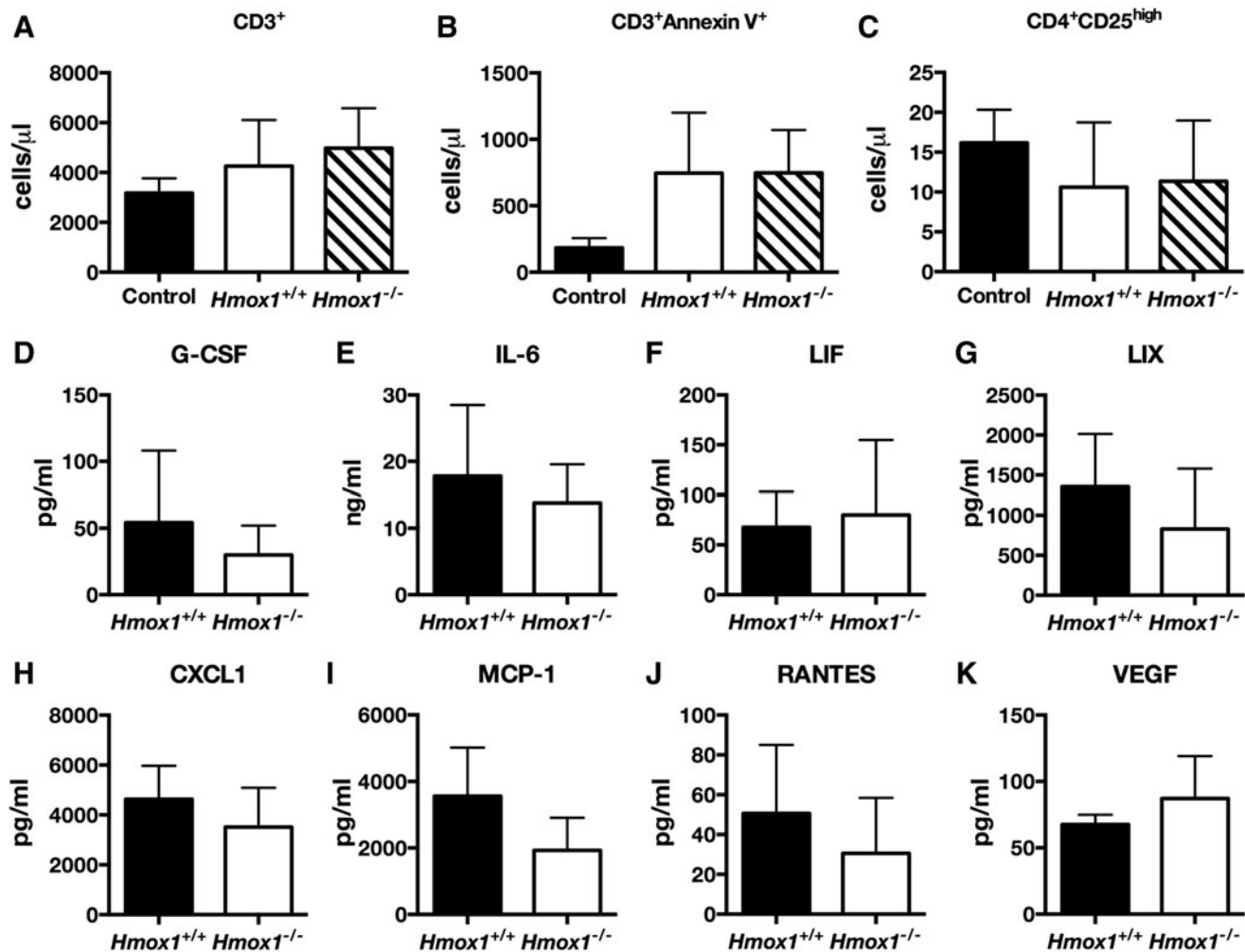


FIG. 3. Lack of *Hmox1* in MSC does not affect their immunomodulatory properties or secretory profile. Number of circulating CD3⁺ T cells (A), CD3⁺AnnexinV⁺ apoptotic T cells (B), or activated CD3⁺CD4⁺CD25^{high} T cells (C) in control mice or injected i.v. with MSC *Hmox1*^{+/+} or *Hmox1*^{-/-} ($N=3-4$). Concentration of G-CSF (D), IL-6 (E), LIF (F), LIX (G), CXCL1 (H), MCP-1 (I), RANTES (J), and VEGF (K) in conditioned media from *Hmox1*^{+/+} or *Hmox1*^{-/-} MSC assessed with multiplex assay on Luminex platform. Data are shown as mean \pm SD, Mann-Whitney, $N=4$.

Subsequently, we checked the toxic concentrations of hemin for MSC *Hmox1*^{+/+} and *Hmox1*^{-/-}. Surprisingly, hemin was more toxic for *Hmox1*^{-/-} MSCs than for *Hmox1*^{+/+} cells not until at a concentration of 200 μ mol/L (Fig. 4C). However, conditioned media from both MSC *Hmox1*^{+/+} and *Hmox1*^{-/-} showed similar total antioxidant capacity assessed with total antioxidant capacity (TAC) (Fig. 4D), 2,2'-azino-bis (3-ethyl-benzothiazoline-6-sulphonic acid) (ABTS), or ferric-reducing antioxidant power (FRAP) assays (Supplementary Fig. S3C, D). The latter result suggests that the reason for the MSC resistance to oxidative stress is rather intrinsic.

First, we supposed that low sensitivity of *Hmox1*^{+/+} and *Hmox1*^{-/-} MSCs to hemin, especially in comparison to BDMC, could be related to the high expression of heme oxygenase-2. However, the expression of *Hmox1* was lower in MSCs *Hmox1*^{+/+} than in PACs *Hmox1*^{+/+} (Fig. 4E). Similarly, the expression of *Hmox2* was lower in MSCs than in PAC cells and similar in *Hmox1*^{+/+} and *Hmox1*^{-/-} cells (Fig. 4F). Then, we hypothesized that low sensitivity of *Hmox1*^{-/-} MSC cells could be caused by the low uptake of heme from the culture medium. Therefore, we analyzed heme

uptake with two methods—direct measurement of heme in cells stimulated for 2 h with 50 μ mol/L hemin, and then after 2 h in fresh medium (Fig. 5A), and with the measurement of tin protoporphyrin IX fluorescence.

Hmox1^{+/+} and *Hmox1*^{-/-} MSCs displayed comparable levels of cellular heme at all timepoints tested (Fig. 5B). Moreover, both cell genotypes showed comparable SnPP fluorescence proportional to the concentration of SnPP in the culture medium (Supplementary Fig. S4A). Therefore, higher resistance to hemin in comparison to PAC cannot be explained by the compensation of *Hmox1* function by *Hmox2* or by changed uptake of hemin.

Since bone marrow PACs are mostly of monocytic origin, and therefore distant from MSCs, we decided to further focus on the differences in stress response between MSCs and fibroblasts. Interestingly, mouse tail-tip fibroblasts had a phenotype similar to MSCs, that is, they expressed Sca1, CD106 and part of them showed positive for CD140a (Supplementary Fig. S5). Both fibroblasts and MSCs, regardless of *Hmox1*, were characterized by the similar expression of heme transporters *Hcp1* (*Slc46a1*) and *Hrg1* (*Slc48a1*)

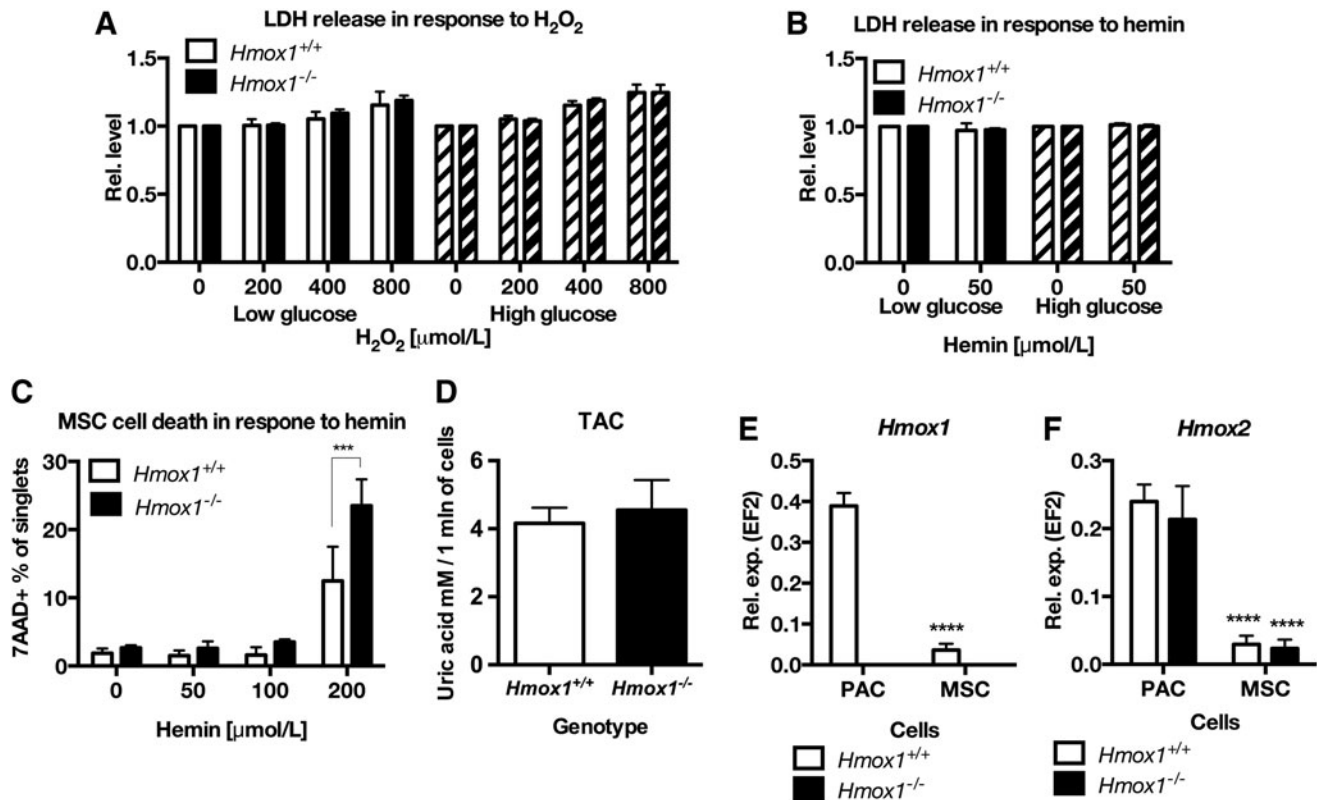


FIG. 4. MSC, regardless of the low expression of both heme oxygenases 1 or 2, show high resistance to hemin or hydrogen peroxide. LDH release in MSC $Hmox1^{+/+}$ or $Hmox1^{-/-}$ treated with H_2O_2 (A) or hemin (B) in low or high glucose medium. Cell death in MSC $Hmox1^{+/+}$ or $Hmox1^{-/-}$ treated with increasing concentrations of hemin for 6 h, assessed with 7-AAD staining and flow cytometry (C). Data are shown as mean \pm SD. *** $p < 0.001$ two-way ANOVA with Bonferroni post-test, $N = 3$. Total antioxidant capacity of conditioned media from MSC $Hmox1^{+/+}$ or $Hmox1^{-/-}$, measured with TAC kit ($N = 4$) (D). (E) Expression of $Hmox1$, (F) $Hmox2$ in murine bone marrow PAC and MSCs isolated from $Hmox1^{+/+}$ or $Hmox1^{-/-}$ mice, **** $p < 0.0001$ MSC versus PAC; two-way ANOVA with Bonferroni post-test, $N = 3$. 7-AAD, 7-aminoactinomycin D; ANOVA, analysis of variance; LDH, lactate dehydrogenase; PAC, pro-angiogenic cell; TAC, total antioxidant capacity.

(Supplementary Fig. S4B, C). MSCs $Hmox1^{-/-}$ were the only ones to increase in response to hemin treatment and the expression of heme exporter *FLVCR1* (Fig. 5C), which may protect them from the heme overload.

To further elucidate the effects of heme on MSCs, we analyzed genes involved in heme synthesis, of which *Alas1* is regulated by heme. Treatment with hemin (50 $\mu\text{mol/L}$) decreased *Alas1* expression in all treated cells. Therefore, we concluded that heme in the culture medium enters treated cells regardless of $Hmox1$ expression and affects known heme-regulated pathways. In $Hmox1^{-/-}$ fibroblasts, already control cells were characterized by lower *Alas1* levels than $Hmox1^{+/+}$ cells (Fig. 5D). Expression of *Uros* in hemin-treated $Hmox1^{-/-}$ fibroblasts was lower than in corresponding MSC cells (Supplementary Fig. S4D).

What is more, $Hmox1^{-/-}$ fibroblasts treated with hemin decreased *Cpox* expression whereas it remained unchanged in other cell types (Supplementary Fig. S4E). There were no differences in the expression of *Hmbs*, *Alad*, *Ppox*, and *Fech* (Supplementary Fig. S4E, F). $Hmox1$ was similarly upregulated in both $Hmox1^{+/+}$ MSCs and fibroblasts (Fig. 5E). The expression of $Hmox2$ was not affected by hemin treatment and did not differ in any of the tested groups of cells (Fig. 5F). To sum up, MSCs $Hmox1^{-/-}$ show higher resistance to H_2O_2

and hemin than previously tested PACs (16). However, low sensitivity to hemin is not caused by changes in heme uptake or synthesis and it is not related to $Hmox2$.

Hemin increases cellular H_2O_2 in $Hmox1^{-/-}$ MSCs and fibroblasts

To assess the effects of hemin on oxidative stress in the MSCs $Hmox1^{+/+}$ and $Hmox1^{-/-}$, we analyzed the levels of cellular hydrogen peroxide by using H_2DCFDA staining. Tail-tip fibroblasts isolated from the same $Hmox1^{+/+}$ or $Hmox1^{-/-}$ mice were used as nonprogenitor internal control cells. After 6 h of incubation with hemin (50 $\mu\text{mol/L}$), levels of H_2O_2 were increased in $Hmox1^{-/-}$ MSCs and fibroblasts and higher than in respective wild-type cells (Fig. 5G). After a 24-h incubation period, levels of H_2O_2 remained low in $Hmox1^{+/+}$ cells and were higher in $Hmox1^{-/-}$ fibroblasts than in $Hmox1^{-/-}$ MSCs (Fig. 5H). This suggests that murine MSCs might be equipped with an anti-oxidant protective mechanism, which works better than in fibroblasts.

Nevertheless, after 48 h of stimulation with hemin, both $Hmox1^{-/-}$ MSCs and $Hmox1^{-/-}$ fibroblasts were dead. Interestingly, at this late timepoint, levels of cellular H_2O_2 were higher in hemin-treated $Hmox1^{+/+}$ fibroblasts than in

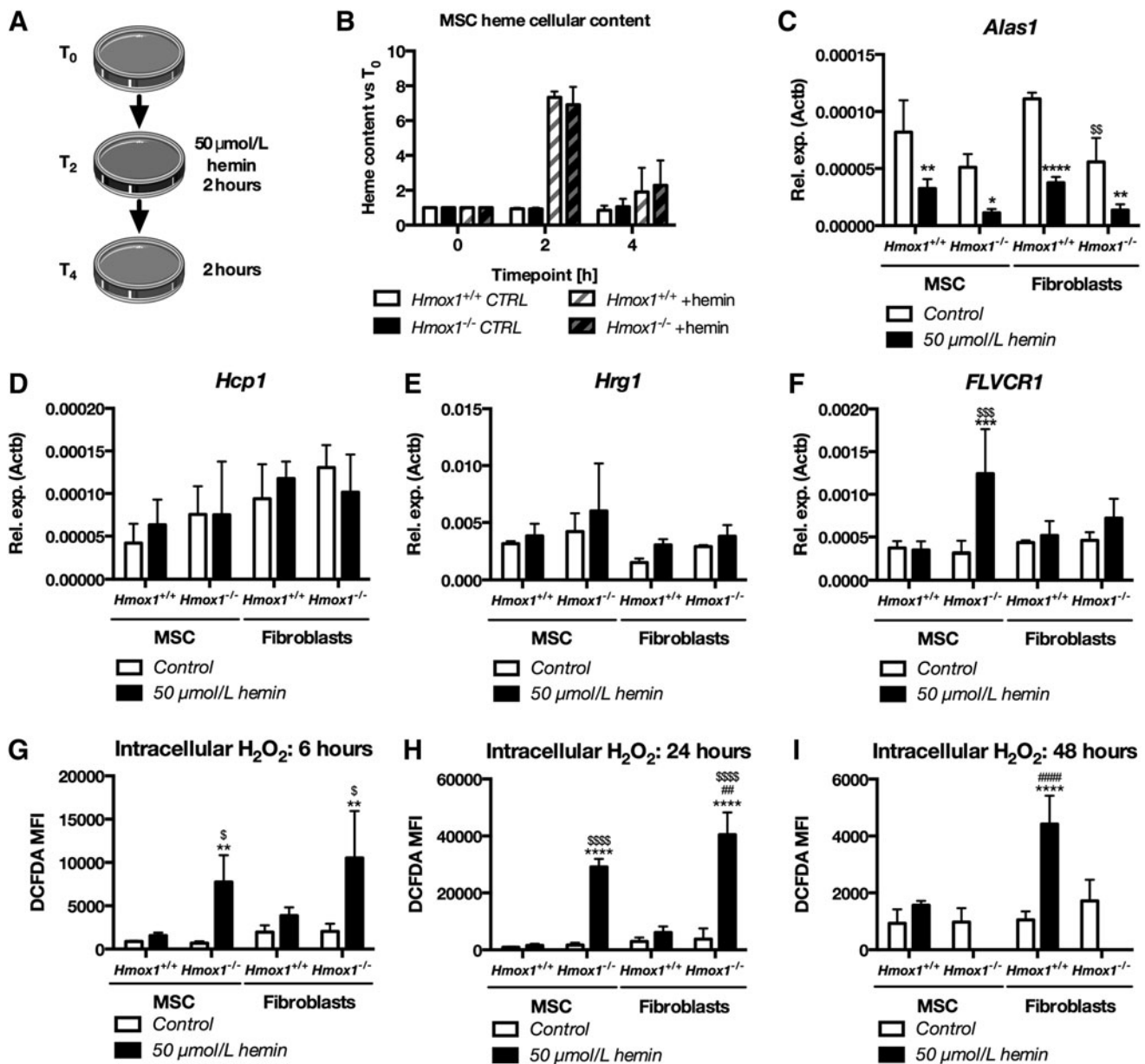


FIG. 5. Hemin increases intracellular hydrogen peroxide in *Hmox1*^{+/+} or *Hmox1*^{-/-} and decreases *Alas1* expression but does not affect levels of heme importers. Scheme of the experiment for the assessment of intracellular heme content (A). Heme cellular content measured with spectrophotometry in MSC *Hmox1*^{+/+} or *Hmox1*^{-/-} stimulated for 2 h with hemin (T₂) and then after 2 h in hemin-free medium (T₄), *N* = 3 (B). Expression of *FLVCR1* (C), *Alas1* (D), *Hmox1* (E), and *Hmox2* (F) in *Hmox1*^{+/+} or *Hmox1*^{-/-} MSCs or fibroblasts stimulated for 6 h with 50 μ mol/L hemin. Levels of H₂O₂ measured with H₂DCFDA after 6 (G), 24 (H), or 48 h (I) in *Hmox1*^{+/+} or *Hmox1*^{-/-} MSCs or fibroblasts stimulated with hemin (50 μ mol/L). Data are shown as mean \pm SD, **p* < 0.05, ***p* < 0.01, ****p* < 0.001, *****p* < 0.0001 hemin-treated cells versus control; ##*p* < 0.01, #####*p* < 0.0001 MSC versus fibroblasts, \$*p* < 0.05, \$\$*p* < 0.01, \$\$\$*p* < 0.001, \$\$\$\$*p* < 0.0001 *Hmox1*^{+/+} versus *Hmox1*^{-/-}. Two-way ANOVA with Bonferroni post-test, *N* = 3.

Hmox1^{+/+} MSCs (Fig. 5I). Therefore, we concluded that both *Hmox1*^{+/+} and *Hmox1*^{-/-} MSC cells showed higher resistance to hemin than respective tail-tip fibroblasts.

MSCs lacking *Hmox1* efficiently induce antioxidant gene response

A lower concentration of H₂O₂ in MSC *Hmox1*^{-/-} than *Hmox1*^{-/-} fibroblasts after short incubation with hemin sug-

gested that MSCs might more effectively respond to the pro-oxidative insult. Therefore, we evaluated the expression of a vast panel of antioxidant genes in MSCs and *Hmox1*^{+/+} or *Hmox1*^{-/-} fibroblasts in response to 50 μ mol/L hemin. Treatment with hemin did not change the expression of the major regulator of antioxidant gene response *Nfe2l2* (Fig. 6A), which encodes for the Nrf2 transcription factor. Levels of *Sqstm1*, which is both target and regulator of Nrf2 (21), were increased in MSCs and *Hmox1*^{-/-} fibroblasts but the upregulation was higher in MSCs (Fig. 6B).

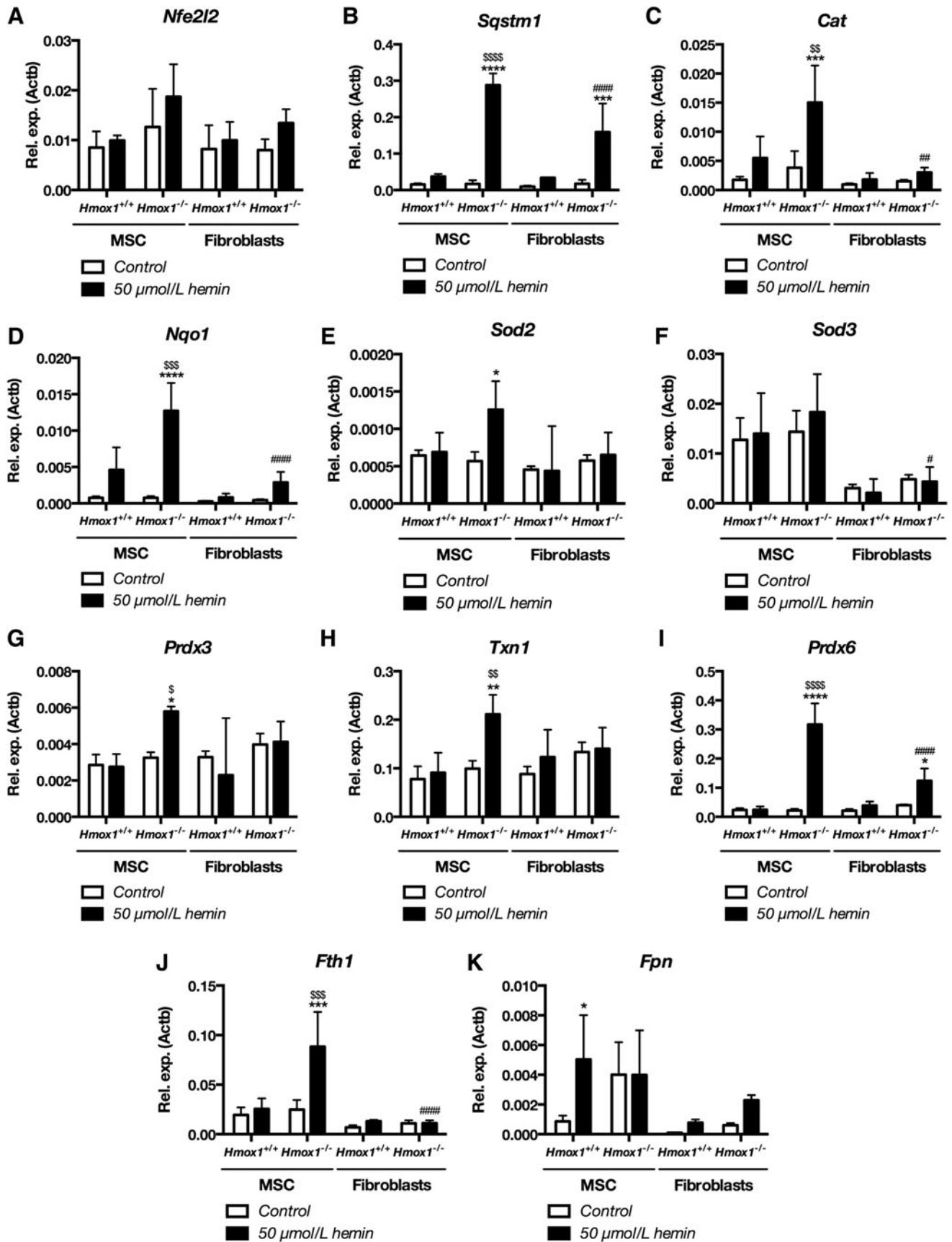


FIG. 6. MSC Hmox1^{+/+} show more efficient antioxidant response after treatment with hemin than fibroblasts Hmox1^{+/+}. Expression of *Nfe2l2* (A), *Sqstm1* (B), *Cat* (C), *Nqo1* (D), *Sod2* (E), *Sod3* (F), *Prdx3* (G), *Txn1* (H), *Prdx6* (I), *Fth1* (J), and *Fpn* (K) in Hmox1^{+/+} or Hmox1^{-/-} MSCs or fibroblasts stimulated for 6 h with hemin (50 μ mol/L). Data are shown as mean \pm SD, * p < 0.05, ** p < 0.01, *** p < 0.001, **** p < 0.0001 hemin-treated cells versus control; # p < 0.05, ## p < 0.01, ### p < 0.001 MSC versus fibroblasts, \$ p < 0.05, \$\$ p < 0.01, \$\$\$ p < 0.001, \$\$\$\$ p < 0.0001 Hmox1^{+/+} versus Hmox1^{-/-}, Two-way ANOVA with Bonferroni post-test, N = 3.

Expression of *Cat* was the highest in hemin-treated MSC *Hmox1*^{-/-}, the only ones to change *Cat* levels (Fig. 6C). Then, *Hmox1*^{-/-} fibroblasts treated with hemin had lower expression of *Nqo1* than *Hmox1*^{-/-} MSCs, the only cells to upregulate *Nqo1* and with its expression higher than *Hmox1*^{+/+} MSCs (Fig. 6D). Similarly, expression of *Sod2* was changed only in *Hmox1*^{-/-} MSC (Fig. 6E) whereas hemin-treated *Hmox1*^{-/-} fibroblasts had lower *Sod3* levels than corresponding MSC cells (Fig. 6F).

Further, expression of *Prdx3* and *Txn1* was enhanced with hemin only in *Hmox1*^{-/-} MSC cells (Fig. 6G, H); whereas *Prdx4*, *Prdx5*, and *Txnrd3* were not affected (Supplementary Fig. S6A–C). Interestingly, *Prdx6*, the only 1-Cys member of peroxiredoxin family (13), was potentially upregulated in hemin-treated *Hmox1*^{-/-} MSCs. *Prdx6* expression was then higher in MSC *Hmox1*^{-/-} than in all other cells (Fig. 6I).

MSCs *Hmox1*^{-/-} were the only ones that upregulated ferritin heavy chain 1 (*Fth1*) expression in response to the hemin treatment (Fig. 6J). Levels of *Fpn* were, on the other hand, increased only in *Hmox1*^{+/+} MSCs (Fig. 6K). Hemin did not affect *Nox4* expression in any cells (Supplementary Fig. S6D). Interestingly, only *Hmox1*^{-/-} MSCs decreased *Gpx1* expression in response to hemin (Supplementary Fig. S6E), whereas expression of *Gpx3* and *Gpx4* was unchanged (Supplementary Fig. S6F, G), and expression of *Gpx8* was decreased in both MSC cell types (Supplementary Fig. S6H). On the other hand, control fibroblasts expressed lower levels of *Gpx8* than respective MSCs (Supplementary Fig. S6H).

To summarize, both MSCs and fibroblasts *Hmox1*^{-/-} upregulated *Sqstm1* and *Prdx6* in response to hemin. Moreover, the expression of both genes was higher in treated MSC *Hmox1*^{-/-} than fibroblasts *Hmox1*^{-/-}. MSC *Hmox1*^{-/-} but not fibroblasts *Hmox1*^{-/-} elevated levels of transcripts for *Cat*, *Nqo1*, *Prdx3*, *Txn1*, and *Fth1*, which confirmed their more efficient antioxidant response.

Importantly, MSC *Hmox1*^{-/-} were the only cells to potentially upregulate quintessential genes involved in the glutathione pathway, namely *Gclc*, *Gclm*, *Gss*, and *Gsr*. MSC *Hmox1*^{-/-} also elevated *Gstp1* (Fig. 7A–E), which forms a heterodimer with *Prdx6* (13), also increased in hemin-treated MSC *Hmox1*^{-/-}. Levels of *Gstk1*, another glutathione S-transferase, remained unchanged in all cell types (Fig. 7F). Levels of the upregulated glutathione pathway genes were higher in hemin-treated *Hmox1*^{-/-} MSC cells than in hemin-treated *Hmox1*^{+/+} MSCs and *Hmox1*^{-/-} fibroblasts. The latter ones increased expression only of *Gclm* and *Gstp1* (Fig. 7B, F).

Finally, to functionally validate the unique influence of hemin on glutathione pathway genes in *Hmox1*^{-/-} MSCs, we analyzed reduced glutathione (GSH) to oxidized glutathione (GSSG) ratio, changes in total GSH and GSSG in MSCs, and fibroblasts of both *Hmox1*^{+/+} and *Hmox1*^{-/-} phenotypes. Cells were treated with hemin (50 μ M/L) for 6 h, like in gene expression experiments. However, we then incubated the cells for two more hours in hemin-free complete medium to allow for GSH recovery. The GSH/GSSG ratio that allows assessing cellular oxidative stress was decreased in both *Hmox1*^{+/+} and *Hmox1*^{-/-} MSCs as well as in *Hmox1*^{-/-} fibroblasts. Further, *Hmox1*^{-/-} fibroblasts were characterized by lower GSH/GSSG ratio than *Hmox1*^{-/-} MSC cells, which additionally had a slightly lower ratio than *Hmox1*^{+/+} MSCs (Fig. 7G).

Noteworthy, *Hmox1*^{-/-} MSCs were the only ones that increased total GSH in response to hemin, whereas they caused a decrease in total GSH in *Hmox1*^{+/+} MSCs (Fig. 7H). Levels of GSSG were increased in *Hmox1*^{+/+} MSCs, *Hmox1*^{-/-} MSCs, and *Hmox1*^{-/-} fibroblasts. They were higher in both *Hmox1*^{-/-} cell types than in their respective *Hmox1*^{+/+} counterparts. However, levels of GSSG increased more in *Hmox1*^{-/-} fibroblasts than in *Hmox1*^{-/-} MSCs (Fig. 7I).

MSCs isolated from *Hmox1*^{-/-} mice showed, in comparison to fibroblasts and PAC, higher resistance to hemin. However, hemin still increased cellular concentration of H₂O₂ in both MSCs and fibroblasts *Hmox1*^{-/-}. We show here for the first time that murine MSCs *Hmox1*^{-/-} could better than mouse tail-tip fibroblasts induce antioxidant gene response, especially genes involved in the glutathione pathway. Importantly, changes in the glutathione pathway, observed on the mRNA level, were further validated functionally by the measurement of reduced and oxidized cellular glutathione. One may speculate that such a fast response to stress factors can, in part, contribute to the presence of only minor effects of *Hmox1* knockout on other functions of murine bone marrow-derived MSCs.

Discussion

Murine bone marrow-derived MSCs isolated from *Hmox1*^{+/+} or *Hmox1*^{-/-} mice showed similar phenotype, proliferation, and differentiation. In our hands, neither proliferation nor differentiation to primary lineages was affected by increased glucose concentrations in the culture media. However, we show here for the first time that in comparison to tail-tip fibroblasts, murine MSCs are more efficient in antioxidant response.

Available reports on the role of *Hmox1* in MSC differentiation are often inconsistent, which may result from the use of cobalt or tin protoporphyrins to modulate *Hmox1* activity or differences between mouse and human MSCs. Such species-dependent variations were reported, for example, for immunomodulatory activities of MSCs, which are regulated by nitric oxide synthase in murine cells and indoleamine-2,3-dioxygenase in human cells (51). We performed our experiments on murine bone marrow-derived MSCs isolated from wild-type or *Hmox1*^{-/-} mice. Importantly, the unaffected phenotype of *Hmox1*^{-/-} MSCs as well as no changes in the differentiation to adipocytes, osteoblasts, and chondrocytes were previously shown by Zarjou *et al.* (66), who also used cells isolated from *Hmox1*^{-/-} mice.

On the other hand, Barbagallo *et al.* reported that expression of heme oxygenase-1 changes during the differentiation of human MSCs to osteoblasts (3) and treatment with osteogenic growth peptide increases *HMOX1* expression in human bone marrow MSCs (56). Moreover, human MSCs stimulated with CoPP during the osteogenic differentiation had upregulated osteonectin, osteogenic growth peptide, and osteocalcin. CoPP decreased adipogenic differentiation of MSCs (3, 56), whereas downregulation of *HMOX1* with siRNA resulted in enhanced adipogenesis.

Human bone marrow MSCs treated with epoxyeicosatrienoic acid displayed decreased levels of Bach1, a repressor of *HMOX1* expression (52), and increased *HMOX1* mRNA; whereas levels of PPAR γ and C/EBP α , involved in adipogenesis, were decreased (57). Vanella *et al.* reported later on

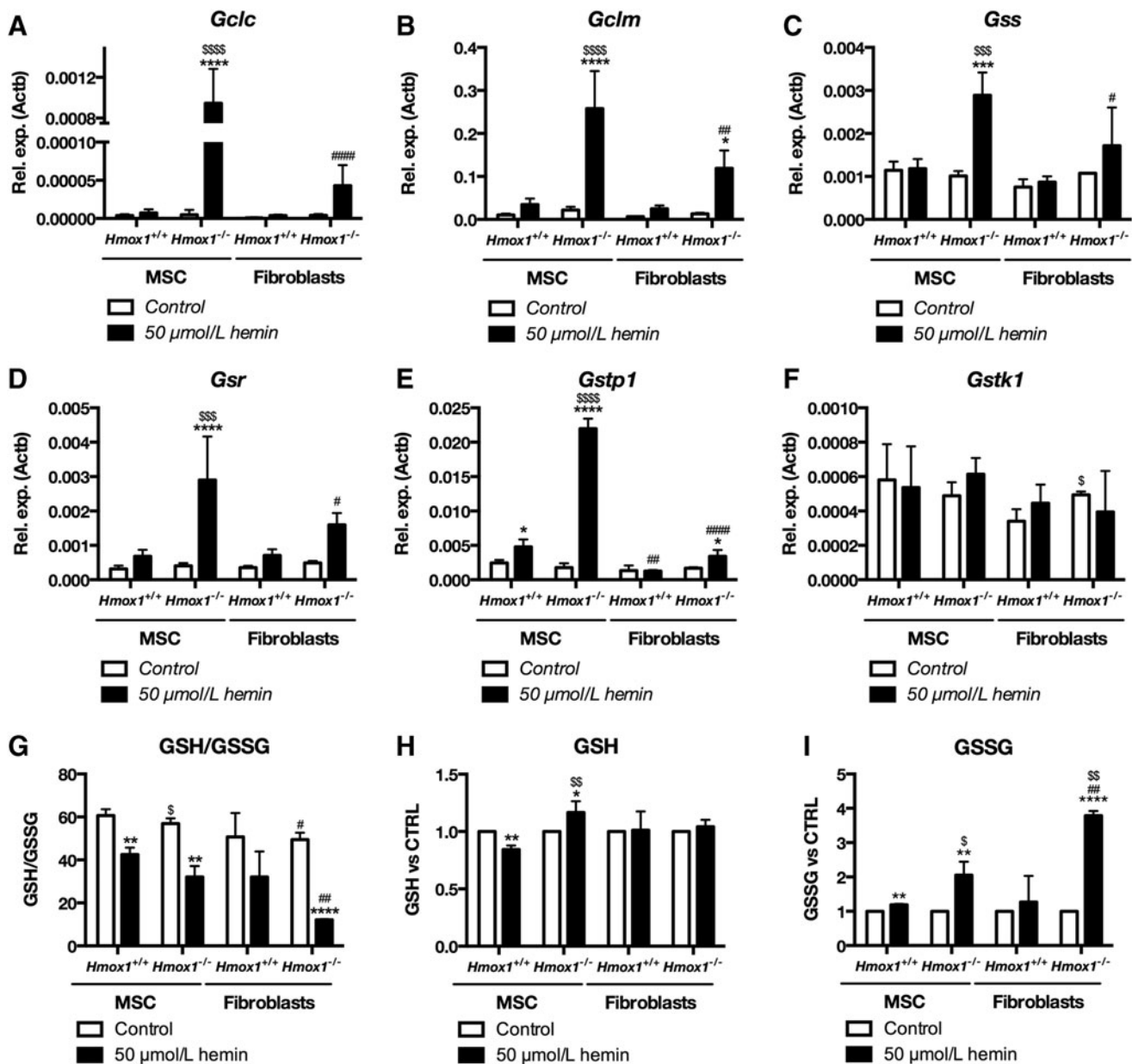


FIG. 7. MSC *Hmxo1*^{+/+} strongly induce expression of glutathione pathway enzymes and increase GSH levels after stimulation with hemin. Expression of *Gclc* (A), *Gclm* (B), *Gss* (C), *Gsr* (D), *Gstp1* (E), and *Gstk1* (F) in *Hmxo1*^{+/+} or *Hmxo1*^{-/-} MSCs or fibroblasts stimulated for 6 h with hemin (50 μmol/L). Ratio of GSH to GSSG (G), changes in total GSH (H), and GSSG (I) in *Hmxo1*^{+/+} or *Hmxo1*^{-/-} MSCs and fibroblasts stimulated for 6 h with hemin (50 μmol/L) and then kept in fresh medium for 2 h. Levels of GSH and GSSG were measured with GSH/GSSG-Glo™ Assay. Data are shown as mean ± SD, **p* < 0.05, ***p* < 0.01, ****p* < 0.001, *****p* < 0.0001 hemin-treated cells versus control; #*p* < 0.05, ##*p* < 0.01, ###*p* < 0.001, \$\$\$*p* < 0.0001 MSC versus fibroblasts, \$*p* < 0.05, \$\$*p* < 0.01, \$\$\$*p* < 0.001, *****p* < 0.0001 *Hmxo1*^{+/+} versus *Hmxo1*^{-/-}, Two-way ANOVA with Bonferroni post-test, *N* = 3. GSH, reduced glutathione; GSSG, oxidized glutathione.

that inhibition of adipogenesis induced with CoPP and increase with tin mesoporphyrin could be linked to modulation of the canonical Wnt signaling (59).

Use of hemin, heme oxygenase-1 substrate, and potent inducer led to the opposite conclusions. Hemin increased adipogenesis in murine 3T3L1 preadipocytes and human bone marrow MSCs but also increased oxidative stress and induced DNA damage in murine preadipocytes (43). Importantly, all observed changes in gene expression induced with hemin could be reversed with antioxidant Tempol. Therefore, the authors concluded that oxidative stress is the

key factor that regulates differentiation patterns in hemin-treated cells (43). Human bone marrow MSCs transduced with adenoviral vectors encoding for *Hmxo1* showed no changes in differentiation pattern but improved viability in hypoxia (18, 68). However, adenoviral vectors give only transient expression of the transgene, which was lost after 2 weeks of culture (18).

Primary rat osteoblasts showed reduced expression and activity of alkaline phosphatase and reduced osteocalcin and Runx2 expression levels when treated with hemin or when transduced with adenoviral vectors harboring the *Hmxo1*

gene (32). Similar results were obtained with carbon monoxide releasing molecule CORM-2 or bilirubin or when *Hmox1* expression was induced with prostaglandin J2 (PGJ2). Effects of hemin or PGJ2 could be reversed with heme oxygenase-1 inhibitor zinc protoporphyrin (32).

In another study, calcification of human smooth muscle cells was inhibited when cells were stimulated with heme (65). Importantly, the effect of heme and heme oxygenase-1 was mediated by ferritin (65). In our study, MSCs isolated from *Hmox1*^{-/-} mice had basal levels of H₂O₂ similar to those of wild-type cells and potentially elevated *Fth1* in response to hemin. Further, *Hmox1*^{-/-} MSCs could induce an efficient antioxidant response to hemin, which is a strong stress factor for cells lacking heme-degrading enzyme. One may speculate that relatively high resistance to oxidative stress observed in MSC *Hmox1*^{-/-} can, at least in part, explain the observed lack of differences in the pattern of differentiation to basic lineages.

MSC *Hmox1*^{+/+} or *Hmox1*^{-/-} tended to induce T cell apoptosis when injected *in vivo*. However, the change did not reach statistical significance. Both previous reports analyzing the role of *Hmox1* in immunosuppressive activity of MSCs used SnPP to inhibit Hmox1 activity (7, 37) and human and rat (7) or human MSCs (37) and did not assess T cell apoptosis *in vivo*. Moreover, an *in vivo* experiment conducted on the role of *Hmox1* in MSCs on the protection from graft rejection lacked the control group of animals treated with SnPP only, without MSCs (7). The role played by *Hmox1* in human cells remained ambiguous, since Mougiakakos *et al.* failed to show the direct relation between *Hmox1* levels and immunomodulatory activity of MSCs (37).

Surprisingly, both *Hmox1*^{+/+} and *Hmox1*^{-/-} MSCs showed high resistance to H₂O₂ or, even more unexpectedly, to hemin, irrespective of the concentration of glucose in the medium. Cells devoid of the heme oxygenase-1, enzyme-degrading heme, were up to now shown to be highly sensitive to hemin (16). Free heme is toxic to the cells and increases oxidative stress that may lead to lipid peroxidation, DNA damage, and protein aggregation [reviewed in (28)].

In our study, we show for the first time that MSCs *Hmox1*^{-/-} were resistant to hemin concentrations, which potentially induced cell death in bone marrow PAC cells (16). Moreover, MSCs *Hmox1*^{-/-} expressed lower basal levels of *Hmox2* than PAC cells, and *Hmox2* remained unaffected by hemin treatment. Subsequently, *Hmox2*-dependent heme degradation cannot be considered a rescue pathway. Finally, *Hmox1*^{+/+} MSCs were characterized by lower *Hmox1* levels than PAC cells and had similar *Hmox1* expression in comparison to fibroblasts.

We hypothesized that high MSC resistance to hemin could result from the low import of free heme. Interestingly, heme uptake in *Hmox1*^{+/+} and *Hmox1*^{-/-} MSCs did not differ. The latter results were mirrored by the expression of *Slc46a1* and *Slc48a1*—heme transporters. Expression of *Slc48a1* was shown, however, to be regulated by Bach1 (61), which represses *Hmox1* and responds to increased heme concentration. In our hands, hemin increased expression of heme exporter *FLVCR1* in *Hmox1*^{-/-} MSCs but not in *Hmox1*^{+/+} MSCs or fibroblasts, regardless of their genotype. Increased *FLVCR1* was previously reported in kidneys of *Hmox1*^{-/-} mice (48). Therefore, we might speculate that an increase in *FLVCR1* can, at least in part, account for the MSC *Hmox1*^{-/-} resistance to hemin.

As expected, hemin decreased expression of 5'-aminolevulinic synthase 1 in all tested cell types. *Alas1* is a heme synthesis rate-limiting enzyme, whose levels are tightly regulated because of the presence of heme regulatory motif in its promoter (38). Other enzymes involved in the heme synthesis were mostly not changed and did not differ between *Hmox1*^{+/+} and *Hmox1*^{-/-} cells. Regarding the iron metabolism, hemin-treated *Hmox1*^{+/+} MSCs upregulated ferroportin. *Fpn* expression did not change in MSC *Hmox1*^{-/-} cells, but they showed a trend toward higher ferroportin levels than wild-type controls. On the other hand, *Hmox1*^{-/-} but not *Hmox1*^{+/+} MSCs or any of fibroblast cells increased ferritin that captures labile iron and, therefore, protects cells from oxidative stress [reviewed in (28)].

Although 6 h of treatment with 50 μ mol/L hemin did not increase cell death in *Hmox1*^{-/-} MSCs, it elevated concentrations of cellular hydrogen peroxide. Levels of H₂O₂ were higher in *Hmox1*^{-/-} fibroblasts than in *Hmox1*^{-/-} MSCs also after 24 h of treatment with hemin and, importantly, in *Hmox1*^{+/+} fibroblasts than in *Hmox1*^{+/+} MSCs after 48 of culture. However, neither in fibroblasts nor in MSCs we found any differences in basal levels of H₂O₂.

Previously, higher H₂O₂ levels were reported in *Hmox1*^{-/-} than in *Hmox1*^{+/+} iPS cells (31). Increased protein carbonylation and lipid peroxidation were also reported in livers and kidneys of *Hmox1*^{-/-} mice (42). Basal intracellular ROS levels in human MSCs and fibroblasts were similar and lower than ROS in INS-1 insulinoma (55). Both MSCs and fibroblasts were characterized by similar levels of *SOD1*, *SOD2*, *CAT*, and *GPX1* mRNA, and higher activities of catalase and glutathione peroxidase-1 than in INS-1 cells (55). Of note, in our study, the concentration of hemin, which was toxic for MSC *Hmox1*^{-/-} cells, also caused some increase in cell death in *Hmox1*^{+/+} MSCs.

MSCs *Hmox1*^{-/-} upregulated in response to hemin a set of genes involved in antioxidant defense, namely *Sod2*, *Prdx3*, *Prdx6*, *Cat*, *Gclc*, *Gclm*, *Gss*, *Gsr*, and *Gstp1*, all of which can be regulated by the Nrf2 transcription factor. Notably, hemin induced expression of enzymes involved in both synthesis and metabolism of glutathione. Bilirubin, which is rapidly formed by biliverdin reductase from biliverdin, a product of heme oxygenase activity, is a strong antioxidant (50) that has properties that are complementary to glutathione (46). Although glutathione has a much higher cellular concentration than bilirubin, it protects mainly hydrophilic proteins. On the other hand, lipophilic bilirubin can protect lipids. However, in our hands, MSC *Hmox1*^{-/-} cells devoid of heme oxygenase-1 did not change the expression of heme oxygenase-2, another source of cellular biliverdin.

Increased expression of γ -glutamylcysteine ligase and glutathione synthetase leads to enhanced production of glutathione, whereas upregulated glutathione reductase restores GSH from GSSG. In our experimental setting, GSH to GSSG ratio, which is an indication of cell redox status, decreased in all tested cells treated with hemin. However, the decrease was much stronger in *Hmox1*^{-/-} fibroblasts than in *Hmox1*^{-/-} MSCs. *Hmox1*^{-/-} fibroblasts were characterized by higher total GSSG levels, whereas *Hmox1*^{-/-} MSCs were the only to increase total GSH.

Of note, HUVEC cells with long alleles of the *HMOX1* promoter, and thus lower levels of HMOX1, treated with H₂O₂ had higher concentrations of total glutathione and GSSG, but

lower GSH/GSSG ratio than cells with the short promoter (53). Further, hemin was shown to induce neuronal necroptosis, which was related to depletion of glutathione (29).

Increased expression of glutathione metabolism genes in *Hmox1*^{-/-} MSCs was accompanied by the upregulation of peroxiredoxin-6. Prdx6 is the only 1-Cys peroxiredoxin that uses glutathione instead of thioredoxin, and works as a heterodimer with glutathione S-transferase π [reviewed in (13)], which was also upregulated in hemin-treated MSC *Hmox1*^{-/-}. Peroxiredoxin-6 has double activity: peroxidase and phospholipase A₂ (13). Interestingly, slightly higher levels of peroxiredoxin 6 were reported in human MSCs than in embryonic stem cells (22) and both peroxiredoxin-6 and glutathione S-transferase π but also peroxiredoxins 1 and 2 were highly abundant in human MSCs (60).

Expression of peroxiredoxin-6 was not changed in late passage MSCs in comparison to early passage MSCs, although aged and more senescent cells showed increased H₂O₂ concentration (19). In another study, aged MSCs were characterized rather by increased peroxiredoxin 5 expression (24). Surprisingly, hemin treatment, which is used to induce erythroid differentiation of K562 erythroleukemia cells, decreased peroxiredoxin-6 levels in K562 cells (25).

We put forward that increased expression of peroxiredoxin 6 and other antioxidant genes should be rather considered as a protective mechanism that allows cells to deal better with oxidative stress than BDMC (Supplementary Fig. S7). Although levels of H₂O₂ in fibroblasts and MSCs were similar, only the latter cells were able to upregulate peroxiredoxin 6 and its partner—glutathione S-transferase π .

MSCs isolated from *Hmox1*^{-/-} mice were able to react to oxidative stress better than fibroblasts and recovered glutathione faster. One may speculate that MSCs can express lower levels of Keap1 or Nrf3, both of which can decrease activity of Nrf2 transcription factor. Nevertheless, our data show that cells such as MSCs are better equipped with the measures to deal with harsh conditions than other bone marrow-derived cells, especially proangiogenic cells. Further, we can speculate that certain cell types are less dependent on heme oxygenase-1, which is considered a crucial cytoprotective enzyme. Tested functions and differentiation potential of murine MSCs were mostly unaffected by the lack of Hmox1.

Materials and Methods

Animals

All procedures involving the use of animals were performed according to approved guidelines. Mice were maintained under the specific pathogen-free conditions, in individually ventilated cages, with full access to food and water. All animal experiments were approved by the Local Ethical Committee for Animal Research at the Jagiellonian University.

Isolation of MSCs

MSCs were isolated from femurs and tibia of C57Bl6 \times FVB *Hmox1*^{+/+} or C57Bl6 \times FVB *Hmox1*^{-/-}. Mice were sacrificed with the overdose of ketamine/xylazine. Bones were resected under the sterile laminar flow hood and cut into small pieces (ca. 1 mm²) with a bone cutter. Then, bone chips were digested

with 1 mg/mL type II collagenase (Gibco) for 90–120 min in 37°C in a rotary shaker (250 rpm). Released cells were washed once with PBS and resuspended in the growth medium [α MEM supplemented with 10% FBS (Lonza) and penicillin with streptomycin (Sigma-Aldrich, St. Louis, MO)].

Cells were seeded in six-well plates—bone marrow from one mouse per well. The medium was changed every 24 h in the first 3 days and every 2–3 days after that, and cells were passaged when confluent. Importantly, before the MACS sorting, cells were detached with short treatment with trypsin (2 min; Gibco) at room temperature to decrease the number of highly adherent macrophages in culture. After three passages, MSCs were further purified from the CD45⁺ fraction with MACS sorting.

MACS sorting of CD45⁻ murine bone marrow stromal cells

Bone marrow-derived cells were detached with trypsin, washed with PBS, resuspended in AutoMACS running buffer (Miltenyi), and stained for 25 min with anti-mouse CD45 MicroBeads (Miltenyi) in 4°C. Then, the cells were washed with 1 mL of PBS, resuspended in AutoMACS Running Buffer, and separated on MACS MS columns (Miltenyi) or with AutoMACS (Miltenyi). Flow-through with CD45⁻ cells was collected, and columns with CD45⁺ cells were discarded. Purified CD45⁻ murine bone marrow MSCs were next counted and either used directly for the experiments or seeded for further culture ($1.5\text{--}2.0 \times 10^4/1\text{ cm}^2$) in α MEM complete medium (CM).

Isolation of murine fibroblasts

Murine adult tail fibroblasts were isolated from C57Bl6 \times FVB *Hmox1*^{+/+} or C57Bl6 \times FVB *Hmox1*^{-/-} according to the previously published protocol (49) and cultured in DMEM (Lonza) medium supplemented with 10% FBS (Lonza) and penicillin with streptomycin (Sigma-Aldrich).

Isolation of murine bone marrow PAC

Murine bone marrow proangiogenic cells were isolated as described earlier (14, 16) and cultured in EGM2-MV medium (Lonza) with 10% FBS and penicillin with streptomycin (Sigma-Aldrich).

Analysis of fibroblast- or osteoblast colony-forming units

Cells isolated from the bone marrow were counted by using Türk solution (Merck) to lyse red blood cells. Then, 1×10^6 of bone marrow cells were seeded per well in six-well plates. Cells were cultured until colonies of fibroblastoid cells were formed. At that stage, part of the wells was fixed and stained with crystal violet and the colonies of fibroblastoid cells were counted. Another part of the cells was treated with osteogenic differentiation medium for the next 3 weeks, and then cells were stained with Alizarin Red S and positive colonies were counted.

MSCs phenotyping

MSCs cultured for three passages after the isolation were detached with trypsin, washed with PBS, and stained for

25 min in AutoMACS Running Buffer at 4°C with the following antibodies: anti-mouse CD45 (clone 30F-11; BD Biosciences), anti-mouse CD29 (clone HM β 1-1; BioLegend), anti-mouse CD31 (clone MEC13.3; BD Biosciences), anti-mouse CD34 (clone RAM34; BD Biosciences), anti-mouse CD90.2 (clone 30-H12; BioLegend), anti-mouse CD105 (clone MJ7/18; BioLegend), anti-mouse CD117 (c-kit) (clone 2B8; eBioscience), anti-mouse CD140a (clone APA5; eBioscience), and anti-mouse Ly-6A/E (Sca-1) (clone D7; eBioscience). The phenotype of the cells was assessed with BD LSR II or BD LSR Fortessa (Becton Dickinson).

MSCs: differentiation to osteoblasts

MSCs *Hmox1*^{+/+} or *Hmox1*^{-/-} were differentiated to osteoblasts with the protocol by Zhu *et al.* (69). Briefly, 2.5×10^4 of sorted CD45⁻ bone marrow stromal cells were seeded per 1 well of 24-well plates. Osteoblast differentiation was induced for 3 weeks with α MEM CM supplemented with 0.1 μ mol/L dexamethasone, 10 mmol/L β -glycerol phosphate, and 50 μ mol/L ascorbate-2-phosphate (all from Sigma-Aldrich) and verified with AlizarinRed S staining (Supplementary Fig. S1A), gene expression analysis, and immunofluorescent staining for osteopontin (clone EPR3688; Abcam). Control cells were cultured in α MEM CM. Low glucose differentiation or control medium contained 5 mmol/L glucose, and high glucose differentiation or control medium contained 33 mmol/L glucose.

MSCs: differentiation to adipocytes

MSCs *Hmox1*^{+/+} or *Hmox1*^{-/-} were differentiated to adipocytes with the protocol by Zhu *et al.* (69). Briefly, 2.5×10^4 of sorted CD45⁻ bone marrow stromal cells were seeded per 1 well of 24-well plates. Adipocyte differentiation was induced for 3 weeks with α MEM CM supplemented with 1.0 μ mol/L dexamethasone, 50 μ mol/L 3-isobutyl-1-methylxanthine (IBMX), and 10 ng/mL insulin (all from Sigma-Aldrich) and verified with OilRedO staining, gene expression analysis, and immunofluorescent staining for Fabp4 (clone EPR3579; Abcam). Control cells were cultured in α MEM CM. Low glucose differentiation or control medium contained 5 mmol/L glucose, and high glucose differentiation or control medium contained 33 mmol/L glucose.

MSCs: differentiation to myofibroblasts

Sorted CD45⁻ bone marrow stromal cells were seeded in 24-well plates (2.5×10^4 /well). Myofibroblast differentiation was induced for 6 days with α MEM CM supplemented with 2 ng/mL recombinant human TGF β 1 (Peprotech) and confirmed with gene expression analysis. Control cells were cultured in α MEM CM.

Analysis of MSC immunosuppressive activity in vivo

C57Bl6 \times FVB mice were injected intravenously with 1 million of MSC *Hmox1*^{+/+} or *Hmox1*^{-/-}, or saline (vehicle control) and sacrificed 24 h later. Numbers of circulating CD3⁺ T cells, CD3⁺AnnexinV⁺ apoptotic T cells, and CD3⁺CD4⁺CD25^{high} activated T cells were assessed by flow cytometry on an LSR Fortessa cytometer (Becton Dickinson). Peripheral blood (PB) was collected in heparinized tubes. Red blood cells were lysed with ammonium chloride red blood cell lysis buffer

(0.15 M NH₄Cl, 10 mM KHCO₃, 0.1 mM EDTA). Obtained total nucleated cells were resuspended in autoMACS Running Buffer (Miltenyi).

Cells were then stained with anti-mouse CD3 (clone 17A2; BD Horizon), anti-mouse CD4 (clone RPA-T4; BD Horizon), anti-mouse CD25 (clone C37; BD Pharmingen), and anti-mouse CD45 (clone 30-F11; BD Pharmingen) antibodies. Annexin V⁺ cells were stained with TACS[®] Annexin V (AnV) kit (Trevigen) according to the manufacturer's instructions. The number of cells per 1 μ L of PB was calculated based on the total leukocyte count (WBC, 10^3 cells/ μ L of PB) and the percentage of each population within the collected events. WBC was measured by using ABC Vet (scil animal care GmbH).

Analysis of MSC immunosuppressive activity in vitro

MSC *Hmox1*^{+/+} or *Hmox1*^{-/-} were seeded on six-well plates. When MSCs reached 100% confluency, they were co-cultured for 24 h with 1 mln of primary mouse splenocytes in each well. After the co-culture, splenocytes were harvested and stained with anti-CD3-AlexaFluor 647 (clone 17A2; BD Pharmingen), anti-Ki67-AlexaFluor 488 (clone B56; BD Pharmingen), and DAPI for the analysis of cell proliferation or with anti-CD3-AlexaFluor 647 (clone 17A2; BD Pharmingen), and Annexin V-FITC (Trevigen) for the analysis of T cell apoptosis. Data are presented as percent of proliferating/apoptotic T cells within the population of CD3⁺ T cells.

Total antioxidant capacity assays

Total antioxidant capacity of conditioned media from MSC *Hmox1*^{+/+} or *Hmox1*^{-/-} was measured with TAC Assay (Cell Biolabs) according to the manufacturer's protocol, or using ABTS, or FRAP method. ABTS assay was performed according to the previously published protocol (40) based on the (12) method. FRAP assay was based on the Benzie and Strain method (4).

Multiplex immunoassays

Levels of factors produced by MSC *Hmox1*^{+/+} or *Hmox1*^{-/-} were measured with Milliplex[®] MAP Mouse Cytokine/Chemokine Bead Panel –32 Plex (Millipore) on Luminex FlexMap 3D platform (Millipore) and analyzed with Milliplex Analyst 3.4 software (Millipore).

Proliferation assay

The proliferation of MSCs was assessed with the BrdU method by using Cell Proliferation ELISA (Roche), according to the manufacturer's protocol. Low glucose medium contained 5 mmol/L glucose, and high glucose medium contained 33 mmol/L glucose. Cells were cultured in high or low glucose medium and BrdU labeling solution for 24 h.

Lactate dehydrogenase activity assay

Cytotoxicity of hemin or H₂O₂ in MSCs was evaluated with CytoTox 96[®] NonRadioactive Cytotoxicity Assay (Promega), according to the manufacturer's protocol. Low glucose medium contained 5 mmol/L glucose, and high glucose medium contained 33 mmol/L glucose. Cells were treated with hemin, H₂O₂, and/or high glucose for 6 h before the analysis.

7-aminoactinomycin D-based cell viability assay

The viability of MSCs stimulated for 6 h with high doses of hemin was assessed by 7-aminoactinomycin D (7-AAD) staining. Stimulated and control cells were detached with trypsin, washed with PBS, and resuspended in AutoMACS Running Buffer (Miltenyi). Then, cells were stained for 10 min with 7-AAD (BD Pharmingen) according to the manufacturer's protocol and analyzed on a BD LSR Fortessa cytometer.

Measurement of MSC viability in response to $TNF\alpha$

MSC $Hmox1^{+/+}$ or $Hmox1^{-/-}$ were incubated with 10 ng/mL $TNF\alpha$ for 24 h. Then, numbers of early and late apoptotic cells were assessed by flow cytometry by using the staining with Hoechst 33342 and 7-AAD according to the protocol by Schmid *et al.* (45).

Measurement of cellular H_2O_2 levels

Levels of cellular H_2O_2 were measured with H_2DCFDA assay. $Hmox1^{+/+}$ or $Hmox1^{-/-}$ MSCs and fibroblasts were stimulated for 6, 24, or 48 h with 50 μ mol/L hemin (Frontiers Scientific). After each timepoint, the stimulated cells and nonstimulated controls were harvested with trypsin, washed with PBS, and stained for 30 min with 0.1 μ mol/L H_2DCFDA (Sigma-Aldrich) in PBS. Then, cells were washed twice with PBS, and DCFDA fluorescence was assessed with a BD LSR Fortessa cytometer (Becton Dickinson).

Heme cellular content assay

MSCs $Hmox1^{+/+}$ or $Hmox1^{-/-}$ were cultured in 24-well plates. Heme content was assessed with the method by Foresti *et al.* (15) in nonstimulated cells (T0), after 2 h of stimulation with 50 μ mol/L hemin (T2) and after 2 additional hours of culture in fresh α MEM CM (T4). Cells were lysed with 80% formic acid (POCH S.A.), and the lysate was transferred to clear plastic 96-well plates. Absorbance was measured at $\lambda = 398$ nm with a GENios microplate reader (Tecan).

SnPP binding assay

Cells stimulated with SnPP (Frontiers Scientific) show fluorescence in APC channel ($\lambda_{ex} = 640$ nm, emission detected with 670/25 bandpass filter). Therefore, we stimulated $Hmox1^{+/+}$ and $Hmox1^{-/-}$ MSCs with 10, 25, or 50 μ mol/L SnPP for 6 h and analyzed their fluorescence with flow cytometry. Cells were detached with trypsin, washed, and resuspended in AutoMACS Running Buffer (Miltenyi). Flow cytometry analysis was performed on a BD LSR Fortessa cytometer (Becton Dickinson).

GSH/GSSG assay

Levels of total GSH, total GSSG, and GSH/GSSG ratio in MSCs and fibroblasts were assessed with GSH/GSSG-Glo™ Assay (Promega) according to the manufacturer's protocol. Then, 1.0×10^4 cells were seeded per well in 96-well plates. Cells were stimulated for 4 h with 50 μ mol/L hemin in α MEM CM and cultured for 2 h in α MEM CM to let them recover GSH levels. Total GSH and total GSSG data are shown as a ratio to the control nonstimulated cells.

Analysis of gene expression

Total RNA was isolated by phenol-chloroform extraction, and it was reverse transcribed with the oligo(dT) primers and RevertAid Reverse transcriptase (Fermentas) or with the NCode™ VILO™ miRNA cDNA synthesis kit (Invitrogen). The expression of genes was assessed by quantitative real-time PCR (qRT-PCR), which was performed in the StepOnePlus system (Applied Biosystems, Foster City, CA) with the specific primers (Supplementary Tables S1 and S2), cDNA and SYBR Green Quantitative RT-PCR kit (Sigma-Aldrich), under conditions summarized in Supplementary Table S3.

Expression of lipid metabolism genes in $Hmox1^{+/+}$ or $Hmox1^{-/-}$ MSCs differentiated to adipocytes was assessed with TaqMan® Array Mouse Lipid-Regulated Genes (Applied Biosystems) and TaqMan Universal PCR Master Mix (Applied Biosystems) with the PCR program described in Supplementary Table S3.

Statistical analysis

Statistical analysis of the data was performed with GraphPad Prism software. Results are expressed as mean \pm SD unless otherwise stated. Statistical significance was accepted at $p < 0.05$. Data obtained in *in vitro* experiments were analyzed with Student's *t*-test when two groups of samples were used. In another case, we used one-way or two-way analysis of variance with Bonferroni post-test. The kind of statistical test applied to analyze given sets of data is provided in the description of figures.

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Author Disclosure Statement

No competing financial interests exist.

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Abbreviations Used

- 7-AAD = 7-aminoactinomycin D
- ABTS = 2,2'-azino-bis (3-ethylbenzothiazoline-6-sulphonic acid)
- BDMC = bone marrow-derived proangiogenic cells
- CM = complete medium
- CoPP = cobalt protoporphyrin IX
- FRAP = ferric-reducing antioxidant power
- GSH = reduced glutathione
- GSSG = oxidized glutathione
- HO-1 = Heme oxygenase-1 (encoded by the HMOX1 gene)
- MSCs = mesenchymal stromal cells
- PAC = proangiogenic cell
- PB = peripheral blood
- PGJ2 = prostaglandin J2
- qRT-PCR = quantitative real-time PCR
- SnPP = tin protoporphyrin IX
- TAC = total antioxidant capacity